WHAT IS CLAIMED IS:

1		1. \ A method for inducing an and	i-tumor response in a	
2	mammalian patient suffering from a tumor comprising			
3	administering to said patient a composition comprising a therapeutically effective			
4	amount of a tumor cell or tumor cell extract that is:			
5	: 1	(i) conjugated to a hapten;		
6	m	(ii) of the same tumor type as the	e patient's tumor;	
7	10	(iii) not allogeneic to said patient	, and	
8	b	(iv) incapable of growing in the b	oody of the patient after	
9	į	injection; and		
10	repeating said administration at weekly intervals.			
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1	sie	2. The method of claim 1, wherein said	d composition is administered	
3	for at least three times.			
1	•	3. The method of claim 1, whe	rein said composition is	
2		administered for at least six times.		
1		4. The method of claim/1 further comp	orising administering a	
2	2 therapeutically effective amount of cyclophosphamide prior to administration of said			
3	3 composition.			
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\mathcal{D}		5. The method of claim 4, wherein cyc	clophosphamide is administered	
(2)	only prior to the first administration of said composition.			
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1		6. The method of claim 4 wherein said	therapeutically effective	
2		amount of cyclophosphamide comprises administering a dose of about 300 mg/M ² of		
3		cyclophosphamide.		
1		7. The method of claim 1 wherein said	d tumor cell or extract is	
2		selected from the group consisting of melanoma, lung, co	olon, breast, kidney, prostate,	
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ovarian and leukemia tumor cell or extract. The method of claim, wherein said tumor cell or extract is a 1 melanoma tumor cell or extract. 2 The method of claim 1 wherein said hapten is selected from the group consisting of dinitrophenyl, trinitrophenyl, N-iodoacetyl-N'-(5-sulfonic 1-naphthyl) ethylene diamine, trinitrobenzenesulfonic acid, fluorescein isothiocyanate, arsenic acid benzene isothiocyanate, trinitrobenzenesulfonic acid, sulfanilic acid, arsanilic acid, dinitrobenzene-S-mustard and combinations thereof. 1 1 1 1 2 2 The method of claim wherein said hapten is dinitrophenyl. Wherein said composition is administered The method of claim 1 2 with an adjuvant. The method of claim 11 wherein said adjuvant is selected from the 12. roup consisting of Bacillus Calmette-Guerin, QS-21, detoxified endotoxin and a cytokine. The method of claim 1 further comprising sensitizing the patient 13. with a therapeutically effective amount of the hapten prior to administering said composition. The method of claim 1 wherein said mammal is not sensitized to 1 14. said hapten prior to administration of said composition. 2 The method of claim 1 wherein said mammal is a human. 15. 1 16. The method of claim 1 wherein said composition comprises a maximum of about 7.5 x 10⁶ tumor cells or c.e. extract per dose.